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PTO/SB/33 (07-09) Approved for use through 07/31/2012. OMB 0651-0031

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PRE-APPEAL BRIEF REQUEST FOR REVIEW		P71362US0		
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I hereby certify that this correspondence is being deposited with the United States Postal Service with sufficient postage as first class mail	Application N	Number	Filed	
in an envelope addressed to "Mail Stop AF, Commissioner for	10/586,99	1	July 21, 2006	
Patents, P.O. Box 1450, Alexandria, VA 22313-1450" [37 CFR 1.8(a)]	,	Culy 21, 2000		
on	First Named	First Named Inventor		
Signature	VORDE	VORDE		
	Art Unit	E	xaminer	
Typed or printed	1793		leng M. Chan	
name			Terig IVI. Chari	
Applicant requests review of the final rejection in the above-identified application. No amendments are being filed with this request. This request is being filed with a notice of appeal.				
The review is requested for the reason(s) stated on the attached sheet(s). Note: No more than five (5) pages may be provided.				
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applicant/inventor.	0/	towly Jas	inature	
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assignee of record of the entire interest. See 37 CFR 3.71. Statement under 37 CFR 3.73(b) is enclosed.	Harvey B. Jacobson, Jr.			
(Form PTO/SB/96)	· · · · ·	Typed o	r printed name	
attorney or agent of record. Registration number. 20,851	202-	202-638-6666		
Registration number 20,65 i			none number	
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attorney or agent acting under 37 CFR 1.34.	Febr	February 28, 2011		
Registration number if acting under 37 CFR 1.34		Date		
NOTE: Signatures of all the inventors or assignees of record of the enti Submit multiple forms if more than one signature is required, see below		ir representative(s) a	re required.	

This collection of information is required by 35 U.S.C. 132. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11, 1.14 and 41.6. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Mail Stop AF, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re A	application of:)	
	Carin Vörde, et al.)	Confirmation No. 9238
Serial	No. 10/586,991)	Art Unit: 1793
Filed:	July 21, 2006)	Examiner: Heng M. Chan
For:	METHOD OF PRODUCING SALTS OF)	February 28, 2011

PRE-APPEAL BRIEF REQUEST FOR REVIEW

MAIL STOP AF

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

This is in response to the Final Rejection mailed August 31, 2010.

Reconsideration of the outstanding rejections is respectfully requested. Claims 1 and 16-34 are pending.

Rejection over Langlet I in view of Latypov

The Examiner has maintained the rejection of claims 1, 18, 23, 28, and 33-34 under 35 U.S.C. § 103(a) as allegedly being unpatentable over Langlet, et al., U.S. Patent No. 5,976,483 ("Langlet I") in view of Latypov, et al., WO 99/46202 ("Latypov"). The Examiner has

recognized that Langlet I differs from the claimed invention in that "it does not teach adding a guanylurea ion to the reaction mixture to form the salt of guanylurea dinitramide." However, the Examiner asserted that it would have been obvious to cure this deficiency because Latypov allegedly teaches that both guanidinium ion (used by Langlet I) and guanylurea ion (used in the presently claimed invention) "are chemical equivalents to make dinitramide salts." In support of this assertion, the Examiner wrote:

"[O]n page 2 of the specification, Applicants recognized that guanidine and gunaylurea are functional equivalents [in] that they both provide desirable positive ions to form sparingly soluble ion pair complexes with dinitramide ion."

(Final Rejection, p. 3). With regards to this sentence, Applicants believe the Examiner has reference to the following sentence in the specification: "The basic nitrogen compound can be, for example, different derivatives of guanidine which form salts which are sparingly soluble in water." However, nothing in this sentence (or anywhere else in the specification) teaches or suggests that guanidinium ion of Langlet I and the guanylurea ion of the present invention are "functional equivalents" or "chemical equivalents" when used in the presently claimed process. The specification merely states that different derivatives of guanidine might form sparingly soluble salts, without identifying any in particular (such as guanylurea ion or guanidinium ion). The specification does not state that guanylurea ion and guanidinium ion are functionally equivalent to each other for use in the claimed invention.

Langlet I discloses a process for producing a salt of dinitramidic acid, which starts with an initial reaction mixture with a very low pH, including a nitrating acid, to form

dinitramidic acid (col. 1, lines 52-66). The dinitramidic acid formed in the first step of Langlet I's process is <u>not</u> stable in the acidic environment and the reaction mixture must be neutralized by the addition of a neutralizing agent to the reaction mixture (see, col. 2, lines 21-23, and, col. 1, lines 46-51). The neutralizing agent is added to interrupt the nitration process at an optimal point (col. 2, lines 23-25).

By contrast, in the presently claimed process only guanylurea is added – without a neutralizing agent - and the final product is directly precipitated from the acidic reaction mixture.

A person of skill in the art seeking to modify Langlet I's process would not use guanylurea ion because guanylurea ion is <u>not</u> a neutralizing agent (i.e. a base). As acknowledged by the Examiner on page 3 of the Final Rejection, guanylurea ion is a weak acid, not a base. By contrast, and contrary to the Examiner's assertion, the guanidinium cation identified by Langlet I as potentially being useful as part of a neutralizing agent - $[C(NH_2)_3]^+$ - is basic (<u>i.e.</u> capable of raising the pH of the reaction solution), having a pKa of about 12.5.

The reaction mixture resulting from the nitration step of Langlet I's process is extremely acidic (pH \leq 0). According to Langlet I, it is necessary to neutralize that solution to produce the desired final product (col. 1, lines 46-51). Therefore, a person of skill in the art would not have sought to substitute guanylurea ion for guanidinium ion in Langlet I's process because guanylurea ion – which is acidic - would not serve to neutralize the extremely acidic solution from the nitration step of Langlet I's process, regardless of the teachings of Latypov.

As pointed out by the Examiner, Langlet I proposes guanidium ion – or guanidinium ion – as a possible neutralizing agent (col. 3, lines 21-22). Both guanidine and guanidium ion (the cationic form of guanidine) form complex slurries of dissolved salts in the Langlet I reaction mixture, which need further processing in order to extract a solid salt (col. 3, lines 31-49). Nothing in Langlet I teaches or suggests that addition of guanylurea ion to the reaction mixture would produce a useful salt directly precipitated from the acidic solution. Thus, a person of skill in the art would have had no reason to try to use guanylurea ion in Langlet I's process. It was surprising, in fact, that guanylurea provides the unusual and useful results that it does (i.e. producing a solid, easily separable precipitate from an acidic mixture).

Finally, Langlet I is crystal clear that in its process, the final dinitramidic salt is precipitated from a neutral reaction mixture. By contrast, in the presently claimed invention, the guanylurea dinitramide is precipitated directly from an acidic solution. Nothing in either Langlet I or Latypov teaches or suggests this step.

Rejection over Langlet I in view of Latypov and Severl

Claims 16-17, 19-22, 24-27, and 29-32 have been rejected under 35 U.S.C. § 103(a) as allegedly unpatentable over Langlet I in view of Latypov and Seyerl, U.S. Patent No. 4,559,409 ("Seyerl"). These claims specifically recite the use of cyanoguanidine in the process of the present invention. As discussed above, the process of the presently claimed invention application would not have been obvious over Langlet I in view of Latypov. Nothing

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in Seyerl remedies these deficiencies. In particular, cyanoguanidine could not function as a

neutralizing agent for the very strongly acidic initial reaction mixture of Langlet I. Thus, a

person of skill in the art would not seek to use cyanoguanidine in Langlet I's process.

In view of the foregoing, Applicants respectfully submit that the rejection over

Langlet I in view of Latypov and Seyerl should be reconsidered and withdrawn.

Conclusion

In view of the foregoing remarks, it is believed that the claims are now in

condition for allowance. If the Examiner believes, for any reason, that personal communication

will expedite prosecution of this application, the Examiner is invited to telephone the

undersigned at the number provided below.

Respectfully submitted,

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